Supplement to:

Clinical and serological features distinguish patients with incomplete lupus classification from systemic lupus erythematosus patients and controls by Aberle, et. al.

Supplementary Table 1. ANA specificities in SLE and ILE patients by indirect immunofluorescence.

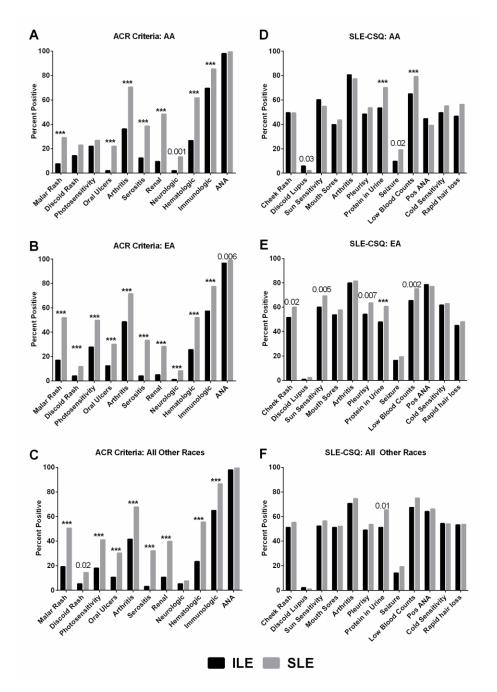
Autoantibody	All SLE	All ILE	OR	
n (%)	(n=3,391)	(n=439)	(95% CI)	Adj. <i>P</i> *
dsDNA	918 (27.0)	40 (9.1)	3.7 (2.6-5.3)	<0.0001
Ro/SSA	716 (21.1)	53 (12.1)	1.9 (1.4-2.7)	<0.001
La/SSB	178 (5.2)	9 (2.1)	2.6 (1.3-5.9)	0.005
Sm	219 (6.5)	7 (1.6)	4.2 (2.0-10.8)	<0.0001
nRNP	684 (20.2)	39 (8.9)	2.6 (1.8-3.7)	<0.0001
ENA-P	52 (1.5)	1 (0.2)	6.8 (1.2-274.3)	0.047+
Jo-1	1 (0.0)	0 (0.0)	NC	1.0

^{*}Bold values are significant (*P*<0.05). *Comparison was made by Fisher's test; all others were performed by Chi-square. NC: not calculable due to 0 value.

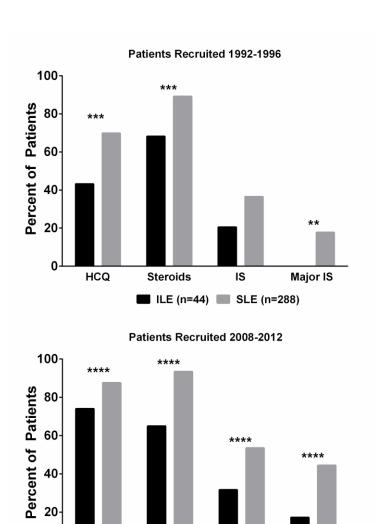
Supplementary Table 2. Autoantibody specificities detected by multiplex assay in African American and European American ILE and SLE patients.

	African American			European American		All Other Races			
	SLE	ILE	Adj.	SLE	ILE	Adj.	SLE	ILE	Adj.
Specificity	(n=978)	(n=103)	P *	(n=1232)	(n=238)	P *	(n=510)	(n=93)	P *
dsDNA, n (%)	350 (35.8)	16 (15.5)	<0.001	287 (23.3)	25 (10.5)	<0.0001	167 (32.7)	7 (7.5)	<0.0001
Chromatin, n (%)	672 (68.7)	33 (32.0)	<0.0001	490 (39.8)	44 (18.5)	<0.0001	283 (55.5)	26 (28.0)	<0.0001
Ribosomal P, n (%)	190 (19.4)	4 (3.9)	<0.001	98 (8.0)	5 (2.1)	0.002	69 (13.5)	2 (2.2)	0.003
Ro/SSA, n (%)	472 (48.3)	27 (26.2)	<0.0001	377 (30.6)	48 (20.2)	0.002	216 (42.4)	33 (35.5)	0.262
La/SSB, n (%)	140 (14.3)	7 (6.8)	0.049	169 (13.7)	20 (8.4)	0.033	83 (16.3)	12 (12.9)	0.505
Sm, n (%)	424 (43.4)	17 (16.5)	<0.0001	176 (14.3)	13 (5.5)	<0.001	130 (25.5)	9 (9.7)	0.001
SmRNP, n (%)	621 (63.5)	39 (37.9)	<0.0001	255 (20.7)	20 (8.4)	<0.0001	191 (37.5)	17 (18.3)	<0.001
RNP, n (%)	555 (56.7)	38 (36.9)	<0.001	250 (20.3)	30 (12.6)	0.007	161 (31.6)	19 (20.4)	0.042
Centromere B, n (%)	33 (3.4)	5 (4.9)	0.399+	58 (4.7)	15 (6.3)	0.382	11 (2.2)	5 (5.4%)	0.084+
Scl-70, n (%)	30 (3.1)	5 (4.9)	0.371+	32 (2.6)	4 (1.7)	0.543	12 (2.4)	2 (2.2%)	1.0+
Jo-1, n (%)	4 (0.4)	2 (1.9)	0.105+	3 (0.2)	0 (0.0)	1.0+	1 (0.2)	0 (0.0%)	1.0+

^{*}Bold values are significant (*P*<0.05). *Comparison was made by Fisher's test; all others were performed by Chi-square.



Supplementary Figure 1. SLE classification criteria and self-reported symptoms in SLE and ILE patients, by race. (A-C) Medical records were reviewed for the 1997 American College of Rheumatology (ACR) SLE classification criteria. (D-F) Self-reported, SLE-related symptoms were determined by patient responses to the SLE-specific portion of the Connective Tissue Disease Screening Questionnaire (SLE-CSQ). Results are shown in African American (A, D), European American (B, E), and other (C, F) SLE and ILE patients. ***P<0.001 by Chisquare or Fisher's test. Exact P values are shown if 0.05>P>0.001. P-values not indicated are not significant (P>0.05).



0

HCQ

Steroids

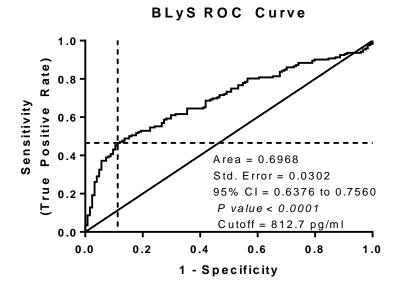
■ ILE (n=174)

IS

SLE (n=1009)

Supplementary Figure 2. Medication usage in SLE and ILE patients during early and late years of recruitment. Medical records were reviewed for use of hydroxychloroquine (HCQ), steroids, immunosuppressants (IS; methotrexate, azathioprine, and sulfasalazine), and major IS (mycophenolate mofetil, cyclophosphamide) for ILE and SLE patients recruited from 1992-1996 (top) or from 2008-2012 (bottom). **P=0.005, ***P<0.001, ****P<0.0001 by Chi-square. P-values not indicated are not significant (P>0.05).

Major IS



Supplementary Figure 3. A cutoff value of 812.7 pg/mL identifies BLyS-positive and BLyS-negative samples. Receiver-operator characteristic (ROC) curves were calculated using quantitative ELISA data for a subset of 72 ILE patients, 100 SLE patients, and 172 controls to determine the BLyS positivity cutoff. The true positive rate (sensitivity) was plotted as a function of the false positive rate (1–specificity) for different positivity thresholds (i.e., different cutoff levels, pg/mL) of a quantitative test. The area under the curve ("Area") is a measure of how well a quantitative test distinguishes between cases and controls.